ANTI-DIARRHEAL- activated attapulgite tablet IBA, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ANTI-DIARRHEAL BOLUS

ADSORBENT, ANTI-DIARRHEAL BOLUS FOR CATTLE

FOR ANIMAL USE ONLY

KEEP OUT OR REACH OF CHILDREN

INDICATIONS

For use as an aid in relief of simple non-infectious diarrhea in horses and cattle.

DOSAGE AND ADMINISTRATION

Administer orally. Give two (2) boluses to adult cattle and horses. Give one (1) bolus to calves and foals.

Repeat treatment at 4 to 6 hour intervals as needed. Discontinue use of product after 3 days. If symptoms persist after using this preparation for 3 days, consult a veterinarian.

COMPOSITION

Each bolus contains: Activated Attapulgite 20%, Carob Flour, Pectin, and Magnesium Trisilicate, in a sugar and starch base.

Store at controlled room temperature between 15°-30°C (59° - 86°F)

TAKE TIME OBSERVE LABEL DIRECTIONS

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DOSAGE AND ADMINISTRATION:

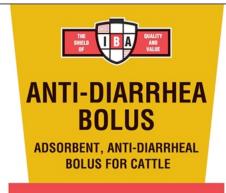
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T-7667-02

Rev. 1-09



CONTENTS: 50 BOLUSES

Manufactured for: IBA Inc. Animal Health Division 27 Providence Road Millbury, MA 01527

COMPOSITION:

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Store at controlled room temperature between 15°-30°C (59°-86°F)

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LOT NO.: EXP. DATE:

IBA STOCK # 474125

99SP2

ANTI-DIARRHEAL

activated attapulgite tablet

Product Type

Product Information

OTC ANIMAL DRUG Item Code (Source) NDC:29983-2709

Route of Administration ORAL

Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
	ATTAPULGITE (UNII: U6 V729 APAM) (ATTAPULGITE - UNII: U6 V729 APAM)	ATTAPULGITE	3.7 g

Product Charac	Product Characteristics			
Color	gray (Light gray to brownish)	Score	2 pieces	
Shape	OVAL (Oblong)	Size	5mm	
Flavor		Imprint Code	2	
Contains				

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:29983-2709-2	50 in 1 JAR		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/28/2007	

Labeler - IBA, Inc. (019494160)

Revised: 3/2018 IBA, Inc.